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FIRST NAMED INVENTOR ATTORNEY DOCKET NO. CONFIRMATION NO. APPLICATION NO. FILING DATE · James E. Brewer A03P1047 4998 10/612,770 07/01/2003 EXAMINER 36802 03/02/2006 7590 PACESETTER, INC. GEDEON, BRIAN T 15900 VALLEY VIEW COURT ART UNIT PAPER NUMBER SYLMAR, CA 91392-9221

3766

DATE MAILED: 03/02/2006

Please find below and/or attached an Office communication concerning this application or proceeding.

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Office Action Summary	Application No.	Applicant(s)	
	10/612,770	BREWER ET AL.	
	Examiner	Art Unit	
	Brian T. Gedeon	3766	
The MAILING DATE of this communication a Period for Reply	ppears on the cover sheet wi	th the correspondence address	
A SHORTENED STATUTORY PERIOD FOR REP WHICHEVER IS LONGER, FROM THE MAILING - Extensions of time may be available under the provisions of 37 CFR after SIX (6) MONTHS from the mailing date of this communication. - If NO period for reply is specified above, the maximum statutory perions are reply within the set or extended period for reply will, by state Any reply received by the Office later than three months after the main earned patent term adjustment. See 37 CFR 1.704(b).	DATE OF THIS COMMUNIC 1.136(a). In no event, however, may a read will apply and will expire SIX (6) MON oute, cause the application to become AB	CATION. eply be timely filed THS from the mailing date of this communication. ANDONED (35 U.S.C. § 133).	
Status			
1) Responsive to communication(s) filed on 01	July 2003.		
2a) This action is FINAL . 2b) ⊠ Th	nis action is non-final.		
3) Since this application is in condition for allow	ance except for formal matt	ers, prosecution as to the merits is	
closed in accordance with the practice under	r Ex parte Quayle, 1935 C.D	. 11, 453 O.G. 213.	
Disposition of Claims			
4)⊠ Claim(s) <u>1-21</u> is/are pending in the application	on.		
4a) Of the above claim(s) is/are withdo			
5) Claim(s) is/are allowed.			
6)⊠ Claim(s) <u>1-7 and 10-20</u> is/are rejected.			
7) Claim(s) 8,9 and 21 is/are objected to.			
8) Claim(s) are subject to restriction and	or election requirement.		
Application Papers			
9) The specification is objected to by the Exami	ner.		
10)⊠ The drawing(s) filed on <u>01 July 2003</u> is/are: a	a)⊠ accepted or b)□ objec	ted to by the Examiner.	
Applicant may not request that any objection to the	ne drawing(s) be held in abeyan	ce. See 37 CFR 1.85(a).	
Replacement drawing sheet(s) including the corre	ection is required if the drawing	s) is objected to. See 37 CFR 1.121(d).	
11) ☐ The oath or declaration is objected to by the	Examiner. Note the attached	Office Action or form PTO-152.	
Priority under 35 U.S.C. § 119			
12) Acknowledgment is made of a claim for foreign a) All b) Some * c) None of: 1. Certified copies of the priority docume 2. Certified copies of the priority docume 3. Copies of the certified copies of the priority docume application from the International Bure	nts have been received. nts have been received in A iority documents have been eau (PCT Rule 17.2(a)).	pplication No received in this National Stage	
* See the attached detailed Office action for a li	st of the certified copies not	received.	
Attachment(s)	" 	(272 4:2)	
1) Motice of References Cited (PTO-892) 2) D Notice of Draftsperson's Patent Drawing Review (PTO-948)		ummary (PTO-413))/Mail Date	
3) ☑ Information Disclosure Statement(s) (PTO-1449 or PTO/SB/0 Paper No(s)/Mail Date <u>7/1/2003</u> .		formal Patent Application (PTO-152)	

DETAILED ACTION

Claim Rejections - 35 USC § 103

The following is a quotation of 35 U.S.C. 103(a) which forms the basis for all obviousness rejections set forth in this Office action:

- (a) A patent may not be obtained though the invention is not identically disclosed or described as set forth in section 102 of this title, if the differences between the subject matter sought to be patented and the prior art are such that the subject matter as a whole would have been obvious at the time the invention was made to a person having ordinary skill in the art to which said subject matter pertains. Patentability shall not be negatived by the manner in which the invention was made.
- 1. Claims 1-7, 10-12 and 14-20 are rejected under 35 U.S.C. 103(a) as being unpatentable over Sloman et al. (US Patent no. 6,738,669) in view of Dahl et al. (US Patent no. 6,976,967).

In regards to claims 1 and 15, Sloman et al. discloses an implantable medical device 10 for multichamber cardiac stimulation that delivers an electrical pulse to one of the chambers of the patient's heart, col 7 lines 47-63. The electrical signals can be delivered to any chamber of the heart, col 11 lines 39-42. Upon delivery of an electrical pulse, far-field sensing occurs, col 11 lines 53-58. For example, if right ventricle stimulation occurs, far-field sensing by a combination of electrodes may occur in the right atrium lead 20 or coronary sinus lead 24, col 12 lines 17-21. Dahl et al. discloses an implantable lead 4 with sensing unit receiving a signal transmitted from a lead disposed in the heart and determines a change in dimension of the heart due to the heart beating, col 6 lines 26-33. Therefore it would have been obvious to one of ordinary skill in the art to use the lead described by Dahl et al. with the methods and apparatus disclosed by Sloman et al. in order to assess the qualitative and quantitative

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aspects of cardiac output relating to the pumping of the heart, capacity, heart rate, and similar cardiac data and better diagnose and treat cardiac symptoms.

In regards to claim 2, Sloman et al. states that the sensing and stimulating electrodes may have a unipolar configuration, col 8 lines 8-12.

In regards to claim 3, Sloman et al. shows that stimulation can take place in the right ventricle, col 12 lines 17-21.

In regards to claim 4, Sloman et al. discloses several leads for placements in the right ventricle with a ring electrode, col 6 lines 47-60.

In regards to claim 5, Sloman et al. teaches that there can be a sensing electrode placed in the superior vena cava, col 6 lines 47-60.

In regards to claim 6, Sloman et al discloses that sensing can occur in a chamber, col 6 lines 47-60, and can occur in any of the four chambers of the heart, col 8 lines 13-17 and col 12 lines 17-21.

In regards to claim 7, Sloman et al. shows that sensing can occur in a unipolar fashion, col 12 lines 17-21.

In regards to claims 10-12, Sloman et al. substantially describes the claimed invention except for the steps of determining a ventricular distance, ventricular volume, or a ventricular distance. Dahl et al. shows that determining the change in dimension of the heart may include a change in the distance between portions of the heart, col 5 lines 40-67. A sensed change in dimension would obviate a calculation of volume since volume is related to spatial dimension. Therefore it would have been obvious to one of ordinary skill in the art at the time the invention was made to use the lead of Dahl et al.

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in combination with the leads of the device and methods of Sloman et al. in order to electrically determine chamber contraction and volumetric output in combination with electrical stimulation therapy.

In regards to claim 14, Sloman et al. states that after a stimulation pulse is delivered to a portion of the heart, the far-field sensing is used in conjunction with other processes to determine if capture in that portion of the heart was successful. If it was determined that capture was unsuccessful, another stimulation pulse is then applied, col 13 lines 26-37.

In regards to claim 16, Sloman et al. substantially describes the claimed invention including a battery for power 110, and stimulation leads 30 having a tip 32 and ring electrode 34, col 6 lines 46-60.

In regards to claim 17, the sensing means of Sloman et al. includes a sensing circuit 82 and 84 coupled to sensing leads 20 and 30, col 8 lines 13-17. Each lead has a plurality of electrodes, col 6 lines 20-60.

In regards to claim 18, the implantable stimulation device 10 has a programmable microcontroller 60 that controls stimulation therapy and processes all incoming signals, col 7 lines 36-47.

In regards to claim 19, Sloman et al. discloses an implantable device 10 with a housing 40, referred to as the "case" or "can", and it may act as the return electrode for all unipolar modes, col 7 lines 5-19. The device has a plurality of leads 20, 24, and 30 connected to the device and is implantable within many structures of the heart, col 6 lines 19-27. Pulse generators 70 and 72 generate stimulation pulses for delivery to the

implantable leads, 20, 24, and 30, col 7 lines 48-63. Sensing circuits 82 and 84 are used upon delivery of an electrical pulse, for far-field sensing occurs, col 8 lines 13-17 and col 11 lines 53-58. For example, if right ventricle stimulation occurs, far-field sensing by a combination of electrodes may occur in the right atrium lead 20 or coronary sinus lead 24, col 12 lines 17-21. Dahl et al. discloses an implantable lead 4 with sensing unit receiving a signal transmitted from a lead disposed in the heart and determines a change in dimension of the heart due to the heart beating, col 6 lines 26-33. Therefore it would have been obvious to one of ordinary skill in the art to use the lead described by Dahl et al. with the methods and apparatus disclosed by Sloman et al. in order to assess the qualitative and quantitative aspects of cardiac output relating to the pumping of the heart, capacity, heart rate, and similar cardiac data and better diagnose and treat cardiac symptoms.

In regards to claim 20, Sloman et al. possesses one or more ventricular leads, col 6 lines 13-18. Lead 24 is configured for the left ventricle, col 6 lines 40-35.

2. Claim 13 is rejected under 35 U.S.C. 103(a) as being unpatentable over Sloman et al. (US Patent no. 6,738,669) in view of Dahl et al. (US Patent no. 6,976,967) and further in view of Mulligan et al. (US Patent no. 6,438,408).

Sloman et al. in view of Dahl et al. substantially describes the claimed invention except for relating the change in cardiac dimensions to congestive heart failure.

Mulligan et al. uses pairs of impedance electrodes 170, 172, 174, 176 to measure the heart chamber volume, col 16 lines 28-49. The methods and apparatus of Mulligan et al. are believed to benefit patient's suffering from heart failure, including congestive

heart failure, col 28 lines 2-4. Therefore it would have been obvious to one of ordinary skill in the art at the time the invention was made to monitor the physical geometry of the heart in order to better assess and electrically treat cardiac abnormalities.

Allowable Subject Matter

3. Claims 8, 9, and 21 are objected to as being dependent upon a rejected base claim, but would be allowable if rewritten in independent form including all of the limitations of the base claim and any intervening claims.

Information Disclosure Statement

4. The information disclosure statement filed 01 July 2003 fails to comply with 37 CFR 1.98(a)(2), which requires a legible copy of each cited foreign patent document; each non-patent literature publication or that portion which caused it to be listed; and all other information or that portion which caused it to be listed. It has been placed in the application file, but the information referred to therein has not been considered. Therefore the Examiner did not consider the cited non-patent literature.

Conclusion

5. The prior art made of record and not relied upon is considered pertinent to applicant's disclosure. Mulligan et al. (US 2005/0027323) discloses an implantable medical device for monitoring cardiac blood pressure and chamber dimension. Ben-Haim et al. (US Patent no. 6,891,091) discloses a method and apparatus for rapidly

generating an electrical map of a chamber of the heart utilizing a catheter. Budd et al.

(US Patent no. 5,662,108) discloses a mapping catheter for mapping a heart chamber

using active electrodes to impose an electrical field within a chamber of the heart, and

passive electrodes to record potentials.

Any inquiry concerning this communication or earlier communications from the

examiner should be directed to Brian T. Gedeon whose telephone number is (571) 272

3447. The examiner can normally be reached on M-F 8:30-5:00.

If attempts to reach the examiner by telephone are unsuccessful, the examiner's

supervisor, Robert E. Pezzuto can be reached on (571) 272 6996. The fax phone

number for the organization where this application or proceeding is assigned is 571-

273-8300.

Information regarding the status of an application may be obtained from the

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Business Center (EBC) at 866-217-9197 (toll-free).

Brian T. Gedeon Patent Examiner Art Unit 3766 Robert E. Pezzuto

Supervisory Patent Examiner

Art Unit 3766

BTG